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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
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10/820,091

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Yulu Wang

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McCARTER & ENGLISH, LLP
Four Stamford Plaza
107 Elm Street
Stamford, CT 06904-0120

EXAMINER

CHANG, ROSIE YUH LOO

ART UNIT

PAPER NUMBER

1762

SHORTENED STATUTORY PERIOD OF RESPONSE	MAIL DATE	DELIVERY MODE
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3 MONTHS

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PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

If NO period for reply is specified above, the maximum statutory period will apply and will expire 6 MONTHS from the mailing date of this communication.

5

Office Action Summary	Application No. 10/820,091	Applicant(s) WANG ET AL.	
	Examiner ROSIE YL CHANG	Art Unit 1762	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 07 April 2004.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-29 is/are pending in the application.
 4a) Of the above claim(s) 29 is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 1-28 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☒ Claim(s) 1-29 are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
 Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
 Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
 a) ☐ All b) ☐ Some * c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
2. ☐ Certified copies of the priority documents have been received in Application No. _____.
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).
- * See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|---|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413)
Paper No(s)/Mail Date. _____ |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | 5) <input type="checkbox"/> Notice of Informal Patent Application |
| 3) <input checked="" type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08)
Paper No(s)/Mail Date <u>See Continuation Sheet</u> . | 6) <input type="checkbox"/> Other: _____ |

Continuation of Attachment(s) 3). Information Disclosure Statement(s) (PTO/SB/08), Paper No(s)/Mail Date :3/28/2005;3/1/2005;2/16/2005;12/10/2004.

DETAILED ACTION

Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121:

- I Claims 1-28, drawn to a method for coating ultra fine particles, classified in class 427, subclass 213.3.
- II Claim 29, drawn to polymer-coated ultra fine particle, classified in class 428, subclass 403.

Inventions I and II are related as process of making and product made. The inventions are distinct if either or both of the following can be shown: (1) that the process as claimed can be used to make another and materially different product or (2) that the product as claimed can be made by another and materially different process (MPEP § 806.05(f)). In the instant case the product as claimed can be made by electrostatic spaying coating with a polymer.

Because these inventions are independent or distinct for the reasons given above and there would be a serious burden on the examiner if restriction is not required because the inventions have acquired a separate status in the art in view of their different classification, restriction for examination purposes as indicated is proper.

During a telephone conversation with Samuel J. DuBoff on 11/20/2006 a provisional election was made with traverse to prosecute the invention of method, claim 1-28. Affirmation of this election must be made by applicant in replying to this Office action. Claim 29 withdrawn from further consideration by the examiner, 37 CFR 1.142(b), as being drawn to a non-elected invention.

Claim Objections

Claim 17 is objected to because of the following informalities: the phrase "temperature is less than the glass transition temperature of the polymer of the polymer." is not proper. Perhaps the phrase "temperature is less than the glass transition temperature of the polymer." would be more clear.

Appropriate correction is required.

Claim Rejections - 35 USC § 112

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Claims 4 and 6 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 4 is rejected, the phrase "the polymer content of said ultra fine particles is from 1 to 100 weight percent based on the total weight of the polymer-coated ultrafine particles" renders the claim indefinite because it is unclear the polymer content of what particle(s) is the claimed invention.

As for claim 6, it is unclear to the Examiner what is the applicant intend to claim in the invention. Is it the "quantity" of the precipitated polymer-coated particles function

to provide controlled release of drug? Or the precipitated polymer-coated particles function to provide controlled release of drug.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

Claims 1-4, 6-11, 18, 20-22, 26-28 are rejected under 35 U.S.C. 102(b) as being anticipated by Subramanian et al. (US 5,833,891).

Subramanian et al. ('891) teach a supercritical fluid process for the coating ultra-fine drug (col. 5, line 6) particle (0.6 micrometer) with polymer (col. 25, claim 21, 22), comprising: preparing a solution of a polymer in an organic solvent, such as acetone (col. 22, line 51), suspending a quantity of ultrafine particles in said solution; and combining a supercritical fluid as antisolvent, such as CO₂, (col. 5, line 30) with said suspension to cause at least a portion of said quantity of suspended ultrafine particles to precipitate from said solution as polymer-coated ultrafine particles (abstract and col. 18, line 65 paragraph of "particle coating "). Subramanian et al. ('891) further teach a variety of core particles can be used in the process (col.7, line 9-15) such as glass or sugar beads or medicament in any discrete solid dosage forms can be coated. This process has many applications in all areas, for example, pharmaceutical application,

pesticide application, polymer application and conductive ink application (col. 22, line 39-44 and col. 23, line 1-8).

As for claim 4:

Subramanian et al. ('891) teach the polymer coating would be from about 1-30% by weight of the final coated product (col.7, line 20-21).

As for claims 10,11,26 and 27:

Subramanian et al. ('891) teach using an apparatus (page 14, Figure 2) for this coating process involving passing the fluid drug-polymer dispersion solution through first passageway into the high-pressure vessel containing the antisolvent (col.5, line 45-59). Simultaneously, an "energizing gas" steam is passed along the second passageway into said vessel. A high-energy sonic wave device is used for break up agglomerates of polymer coated drug particles (col. 5, 60-67).

As for claims 18, 20:

Subramanian et al. ('891) further teach the "energizing gas" is the same as the selected antisolvent, and in most cases carbon dioxide is used both as the antisolvent and "energizing gas" which may be selected from the group consisting of nitrogen, carbon dioxide, propane, trifluoromethane and mixtures thereof (col. 6, line 31-43). As disclosed in the above the same supercritical fluid been used as the antisolvent.

As for claims 7, 8 and 22:

Subramanian et al. ('891) further teach (col. 5, line 5-12) the method involving contacting the drug-polymer dispersion solution with antisolvent at its supercritical fluid condition and cause the antisolvent to deplete the dispersant and precipitate the

Art Unit: 1762

substance as fine particles. Condition enhances the mass transfer rate between the antisolvent and the dispersant so that particle nucleation and precipitation occur rapidly. Then, (Col. 10, line 13-19) the precipitated polymer-coated drug particles are purged with supercritical fluid until the organic solvent is completely depleted from the system (col. 7, line 43-46). The antisolvent is bone dry CO₂ (99.8% purity) (col.13, line 10).

A person shall be entitled to a patent unless –

(e) the invention was described in (1) an application for patent, published under section 122(b), by another filed in the United States before the invention by the applicant for patent or (2) a patent granted on an application for patent by another filed in the United States before the invention by the applicant for patent, except that an international application filed under the treaty defined in section 351(a) shall have the effects for purposes of this subsection of an application filed in the United States only if the international application designated the United States and was published under Article 21(2) of such treaty in the English language.

Claims 1, 2, 5, 6, 9, 18, 21, 28 are rejected under 35 U.S.C. 102(e) as being anticipated by Bertuccio et al. (Drugs encapsulation using a compressed gas antisolvent technique, E. Reverchon (Ed.), Sept.7-10, Capri, Italy, 1997, 327-334).

Bertuccio et al. teach a method of coating ultra fine particles with a polymer (abstract), comprising: preparing a solution of an polymer in an organic solvent; suspending a quantity of ultrafine particles in said solution (page 328, line 19-20); and combining a supercritical fluid as an antisolvent with said suspension to cause at least a portion of said quantity of suspended ultrafine particles to precipitate from said solution as polymer-coated particles. (Page 330, Results and Discussion).

As for claims 2 and 28:

Bertuccio et al. teach the ultra fine particles are pharmaceutical compounds (page 328, Material and Methods) and the supercritical fluid is carbon dioxide (page 328, line 19). Bertuccio et al. teach a method of using the supercritical fluid as the antisolvent to encapsulate a fine drug particle to control drug release rate.

As for claim 5:

Bertuccio et al. teach the polymer used for drug encapsulation is an acrylic polymer, Eudragit E 100, a copolymer of dimethylaminoethyl methacrylate and methacrylic acid ester (page 328, Material and Methods).

As for claim 6:

Bertuccio et al. teach using a model system for the drug (page 328, Material and Methods), spherical particles of potassium chloride, and the effectiveness of drug encapsulation being checked by measuring the conductivity of the encapsulated KCl salt in water. The result is reported in Fig. 5 on page 333. It can be seen in this resolution test, the percentage of "released drug" increasing with time; about 65% of drug released at 10 min. and 95% of drug released at 60 min. Bertuccio et al. teach the polymer-coated ultra fine particles function to provide controlled release of drug.

As for claim 9:

Bertuccio et al. teach the particles used in drug encapsulation are completely soluble in water but totally not soluble in all the organic solvents used to dissolve the polymers (page 328, Materials and Methods).

As for claim 21:

Bertuccio et al. further teach the organic solvent used to dissolve polymer is acetone (page 331, Table 1).

Claims 1,3,5-8,10,11, 14-17, 18 and 26 are rejected under 35 U.S.C. 102(e) as being anticipated by Gupta et al. (US 6,620,351).

Gupta et al ('351) teach a method (col. 6, line 35-55) of using a supercritical fluid as the antisolvent to encapsulate ultra fine particles with polymer (col. 14, line 64-66) to form composite nanoparticles, comprising preparing a solution of polymer, poly (lactide-co-glycolide) in dichloromethane; suspending magnetite nanoparticles (particle size is about 10 nanometer) in said solution (col. 15, line 4-6). The apparatus used in the teaching of Gupta et al. ('351) consisting a particle-polymer dispersion solution delivery system, an antisolvent supply system and a particle production high-pressure vessel (sheet1 of 25, Fig. 1). The particle-polymer dispersion solution is pumped to the high-pressure vessel through a nozzle at a predetermined flow rate (page 9, line 28-41) and the antisolvent, CO₂, is delivered to the bottom of this high-pressure vessel through another pump. The high-pressure vessel is filled with the antisolvent up to the desired operating pressure (page 9, line 53-60) and combining the particle-polymer dispersion with the antisolvent cause at least a portion of suspended ultrafine particles to precipitate from said solution as polymer-coated ultrafine particles. Gupta et al. ('351) further teach this apparatus is equipped with an energy source to cause the vibration (col.9, line 41-44) on a surface within the particle production vessel, and a vibration

Art Unit: 1762

force can be applied to the particle-polymer dispersion solution to break up agglomerates of ultrafine particles formed.

In one embodiment, Gupta et al ('351) teach this encapsulation of one substance can be achieved using another substance to form coated nanoparticles (col.6, line35-37). The core particle to be coated is dispersed in a suitable medium and mixed a polymer solution and sprayed on to the deflecting surface in the high-pressured vessel to obtain very fine polymer-coated drug particles (col. 6, line 54-43). The produced polymer-coated drug particle can be used for controlled release.

As for claims 7 and 8:

Gupta et al. ('351) teach (col.10, line 24-26) the flow rate of CO₂ is maintained high enough so that all the solvents in the dispersion are removed to obtain dry particles.

As for claims 14-17:

Gupta et al. ('351) teach the size (col. 6, line 17-20) of the ultrafine particle depends on the process parameters, such as pressure and temperature of the antisolvent and concentration and flow rate of the polymer solution.

Gupta et al. ('351) teach encapsulating drug particle with poly (lactide-co-glycolide), PLGA (T_g= 40-55C), with antisolvent, CO₂, at pressure of 96.5 bar (9.68 MPa) and at temperature of 35 C, which is below the glass transition temperature (col.14, line 60-64) of the polymer therein. Examiner notice this is the similar pressure, 8.96 MPa, which was chosen in the instance application on page 17, [0185] for minimizing agglomeration of coated particles.

Claims 1, 2, 11, 12, 13, 18, 22 are rejected under 35 U.S.C. 102(e) as being anticipated by Perrut (US 2003/0,031,784).

Perrut ('784) teaches a method of encapsulating very fine particles (page 1, [0012]) with a coating agent, i.e. a polymer, ethylcellulose (page 4, [0054]) by a method employing a supercritical pressure. Perrut ('784) teaches dissolving ethylcellulose in ethyl acetate and mixing amoxicilline particles into the polymer solution, then combining a supercritical fluid, CO₂ with the solution to cause at least a portion of suspended particles to precipitate from solution as polymer-coated particles.

As for claims 2 and 18:

The fine particle is amoxicillin, and the supercritical fluid is carbon dioxide (page 4, [0054]).

As for claim 12:

Perrut ('784) teaches the concentration of the coating agent in the solvent will preferably sufficiently low to avoid a precipitation giving rise to the formation of agglomerates.

As for claims 11, 13:

Perrut ('784) teaches the 4.5% of ethylcellulose in ethyl acetate in the process (page 4, [0054]), the density of ethyl acetate is about 0.897 g/ml at room temperature. Therefore, 4.5% is about 4 mg/ml of polymer in solvent.

As for claim 22

Perrut ('784) teaches (page 3, [0034]) the supercritical fluid percolating within the polymer solution and cause the organic solvent loses a large part of its solvent power.

Art Unit: 1762

Perrut ('784) further teaches (page 3, [0037]) due to the percolation of the supercritical fluid in the solution, the polymer passes into oversaturation and consequently, precipitates on the particles to coat them.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Claim 19 is rejected under 35 U.S.C. 103(a) as being unpatentable over Subramanian et al. (US 5,833,891) in view of Lee (US 6,596,206)

Subramanian et al. ('891) teach that which is disclosed in the above.

Subramanian et al. ('891) teach use supercritical carbon dioxide as the antisolvent and fail to teach using supercritical ammonia or a combination of supercritical fluids as the antisolvent in the SAS process of encapsulation of drug ultrafine particles.

Lee ('206) teaches a SAS process to generate ultra fine particles (about 5 nm to 2.5 um in diameter) of a pharmaceutical agent (col. 7 line 9-47), wherein the solvent is an aqueous or an organic solvent, such as acetone (col.8, line 65), and the antisolvent is a supercritical fluid (col. 16, claim1). Lee (' 209) further teaches (col.5, line 2-7) when the antisolvent is admixed with the solution of drug in solvent; the solubility of the drug

can be reduced to the point at which it precipitates out of solution. The antisolvent is supercritical fluid (col. 10, line 1-15) selected from the group consisting of carbon dioxide, ammonia and combinations thereof (col.16, line 43-54).

Both Subramanian et al ('891) and Lee ('206) utilize supercritical fluid as antisolvent to prepare fine particles of polymer and drug. Supercritical fluids such as ammonia, carbon dioxide and combination thereof are commonly utilized in the art as evidenced by Lee ('206). One having ordinary skill in the art would recognize that choice of one or the other supercritical fluid is depended on organic solvent, which is used to dissolve polymer, solubility in the supercritical fluid. The antisolvent must be at least partially miscible with the organic solvent such the antisolvent is capable of penetrating into polymer solution to cause the desired precipitation of the polymer. Since both Lee ('206) and Subramanian et al ('891) utilize acetone as the organic solvent, therefore, it would have been obvious to one having ordinary skill in the art to utilize supercritical ammonia or a combination of supercritical fluids as taught by Lee ('206) in the method of Subramanian et al ('891) to produce polymer coated drug particles as the supercritical ammonia is easily available and might cost less too.

Claims 23-25 are rejected under 35 U.S.C. 103(a) as being unpatentable over Perrut ('784) in view of Lee ('206).

Perrut ('784) teaches that which is disclosed in the above. Perrut ('784) teaches encapsulation fine particles of amoxilline with ethylcellulose in a SAS process. Perrut

Art Unit: 1762

('784) is silent about formulate the drug particles with diluent or fill such as lactose, dextrose, cellulose and combinations thereof.

Lee ('206) teaches that which is disclosed in the above. Lee ('206) further teach the pharmaceutical agents that are known candidates for administration using dry powder inhalation therapy include, peptidyl drugs and analgesics (Col. 7, line 9-11). Pharmaceutical formulation typically includes components other than the active agent (Col.7, line 51-53) such as a carrier, dextrose, sucrose and fructose (col. 7, line 55-60). Other additives commonly included in a particulate pharmaceuticals included diluents, stabilizers and lubricants (col. 8, line 1-4). For controlled release particles, a biodegradable polymer, such as a polyethylene glycol (page 8, line 29), may be incorporated into the solid particles prepared according to the disclosure above. The weight ratio of the diluents can vary from about 0.1 to 1 about 100,000 to 1 depending upon the application (col.8, line 12-14).

Lee ('491) teaches a SAS process for producing a formulated drug particle not specifically for coating a fine solid drug particle. However, it will be readily apparent to persons skilled in the art, the SAS process for particle formation and for particle coating are similar, except that in coating applications the host particles are suspended in the polymer solution before being delivered into SC CO₂. It would have been obvious to one having ordinary skill in the art to use the formulated drug particles consisting of filler and diluents of the teaching by Lee ('491) in the method of Subramanian et al. ('891) to produce polymer encapsulated drug particles in a similar SAS mass transfer process.

Allowable Subject Matter

There is no allowable subject matter at this time.

Conclusion


The prior art made of record and not relied upon is considered pertinent to applicant's disclosure. Document number A to D and U listed on Notice of References Cited.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to ROSIE YL CHANG whose telephone number is 571-272-6466. The examiner can normally be reached on MONDAY TO FRIDAY 7: 00AM TO 3:30 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, TIMOTHY MEEKS can be reached on 571-272-1423. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Art Unit: 1762

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.



TIMOTHY MEEKS
SUPERVISORY PATENT EXAMINER